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(54) **COATED IMPLANTABLE MEDICAL DEVICE**

BESCHICHTETE, IMPLANTIERBARE MEDIZINISCHE GERÄTE

DISPOSITIF MEDICAL IMPLANTABLE DOTE D'UN REVETEMENT

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EP 0 968 013 B1

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Description

Technical Field

[0001] This invention relates generally to human and veterinary medical devices, and more particularly to devices incorporating drugs or bioactive agents.

Background of the Invention

[0002] It has become common to treat a variety of medical conditions by introducing an implantable medical device partly or completely into the esophagus, trachea, colon, biliary tract, urinary tract, vascular system or other location within a human or veterinary patient. For example, many treatments of the vascular system entail the introduction of a device such as a stent, a catheter, a balloon, a wire guide, a cannula, or the like. However, when such a device is introduced into and manipulated through the vascular system, the blood vessel walls can be disturbed or injured. Clot formation or thrombosis often results at the injured site, causing stenosis or occlusion of the blood vessel. Moreover, if the medical device is left within the patient for an extended period of time, thrombus often forms on the device itself, again causing stenosis or occlusion. As a result, the patient is placed at risk of a variety of complications, including heart attack, pulmonary embolism, and stroke. Thus, the use of such a medical device can entail the risk of precisely the problems that its use was intended to ameliorate.

[0003] Another way in which blood vessels undergo stenosis is through disease. Probably the most common disease causing stenosis of blood vessels is atherosclerosis. Atherosclerosis is a condition which commonly affects the coronary arteries, the aorta, the iliofemoral arteries and the carotid arteries. Atherosclerotic plaques of lipids, fibroblasts, and fibrin proliferate and cause obstruction of an artery or arteries. As the obstruction increases, a critical level of stenosis is reached, to the point where the flow of blood past the obstruction is insufficient to meet the metabolic needs of the tissue distal to (downstream of) the obstruction. The result is ischemia.

[0004] Many medical devices and therapeutic methods are known for the treatment of atherosclerotic disease. One particularly useful therapy for certain atherosclerotic lesions is percutaneous transluminal angioplasty (PTA). During PTA, a balloon-tipped catheter is inserted in a patient's artery, the balloon being deflated. The tip of the catheter is advanced to the site of the atherosclerotic plaque to be dilated. The balloon is placed within or across the stenotic segment of the artery, and then inflated. Inflation of the balloon "cracks" the atherosclerotic plaque and expands the vessel, thereby relieving the stenosis, at least in part.

[0005] While PTA presently enjoys wide use, it suffers from two major problems. First, the blood vessel may

suffer acute occlusion immediately after or within the initial hours after the dilation procedure. Such occlusion is referred to as "abrupt closure." Abrupt closure occurs in perhaps five percent or so of the cases in which PTA is employed, and can result in myocardial infarction and death if blood flow is not restored promptly. The primary mechanisms of abrupt closures are believed to be elastic recoil, arterial dissection and/or thrombosis. It has been postulated that the delivery of an appropriate agent (such as an antithrombic) directly into the arterial wall at the time of angioplasty could reduce the incidence of thrombotic acute closure, but the results of attempts to do so have been mixed.

[0006] A second major problem encountered in PTA is the re-narrowing of an artery after an initially successful angioplasty. This re-narrowing is referred to as "restenosis" and typically occurs within the first six months after angioplasty. Restenosis is believed to arise through the proliferation and migration of cellular components from the arterial wall, as well as through geometric changes in the arterial wall referred to as "remodeling." It has similarly been postulated that the delivery of appropriate agents directly into the arterial wall could interrupt the cellular and/or remodeling events leading to restenosis. However, like the attempts to prevent thrombotic acute closure, the results of attempts to prevent restenosis in this manner have been mixed.

[0007] Non-atherosclerotic vascular stenosis may also be treated by PTA. For example, Takayasu arteritis or neurofibromatosis may cause stenosis by fibrotic thickening of the arterial wall. Restenosis of these lesions occurs at a high rate following angioplasty, however, due to the fibrotic nature of the diseases. Medical therapies to treat or obviate them have been similarly disappointing.

[0008] A device such as an intravascular stent can be a useful adjunct to PTA, particularly in the case of either acute or threatened closure after angioplasty. The stent is placed in the dilated segment of the artery to mechanically prevent abrupt closure and restenosis. Unfortunately, even when the implantation of the stent is accompanied by aggressive and precise antiplatelet and anticoagulation therapy (typically by systemic administration), the incidence of thrombotic vessel closure or other thrombotic complication remains significant, and the prevention of restenosis is not as successful as desired. Furthermore, an undesirable side effect of the systemic antiplatelet and anticoagulation therapy is an increased incidence of bleeding complications, most often at the percutaneous entry site.

[0009] Other conditions and diseases are treatable with stents, catheters, cannulae and other devices inserted into the esophagus, trachea, colon, biliary tract, urinary tract and other locations in the body, or with orthopedic devices, implants, or replacements. It would be desirable to develop devices and methods for reliably delivering suitable agents, drugs or bioactive materials directly into a body portion during or following a medical

procedure, so as to treat or prevent such conditions and diseases, for example, to prevent abrupt closure and/or restenosis of a body portion such as a passage, lumen or blood vessel. As a particular example, it would be desirable to have devices and methods which can deliver an antithrombic or other medication to the region of a blood vessel which has been treated by PTA, or by another interventional technique such as atherectomy, laser ablation, or the like. It would also be desirable that such devices would deliver their agents over both the short term (that is, the initial hours and days after treatment) and the long term (the weeks and months after treatment). It would also be desirable to provide precise control over the delivery rate for the agents, drugs or bioactive materials, and to limit systemic exposure to them. This would be particularly advantageous in therapies involving the delivery of a chemotherapeutic agent to a particular organ or site through an intravenous catheter (which itself has the advantage of reducing the amount of agent needed for successful treatment), by preventing stenosis both along the catheter and at the catheter tip. A wide variety of other therapies could be similarly improved. Of course, it would also be desirable to avoid degradation of the agent, drug or bioactive material during its incorporation on or into any such device.

Summary of the Invention

[0010] The foregoing problems are solved and a technical advance is achieved in an illustrative vascular stent or other implantable medical device that provides a controlled release of an agent, drug or bioactive material into the vascular or other system, or other location in the body, in which a stent or other device is positioned. Applicants have discovered that the degradation of the agent, a drug or a bioactive material that is applied to such a device can be avoided by positing a coating layer on one surface of the device structure. The agent, drug or bioactive material is posited over at least a portion of the coating layer, wherein the coating layer provides for a controlled release of the bioactive material posited thereon.

[0011] In one aspect, the coating layer comprises a non-porous material of for example a parylene derivative. This coating layer has a thickness preferably in a range from 50 to 500,000 Angstroms, more preferably in a range from 100,000 to 500,000 Angstroms, and illustratively approximately 200,000 Angstroms. In another aspect, the non-porous material is either an adsorbent or an absorbent material, where the coating layer of the adsorbent material has a thickness of approximately 230,000 Angstroms.

[0012] In another aspect of the invention, the bioactive material layer includes a chimeric monoclonal antibody such as an antiplatelet GP IIb/IIIa antibody.

[0013] In still another aspect of the invention, an adhesive promotion layer is posited on one surface of the structure on which the coating layer is posited over at

least a portion thereof. Preferably the adhesion promotion layer includes silane having a thickness in range of 0.5 to 5,000 Angstroms.

[0014] Preferably, when the device is intended for use in the vascular system, the bioactive material in the at least one layer is heparin or another antiplatelet or antithrombotic agent, or dexamethasone, dexamethasone acetate, dexamethasone sodium phosphate, or another dexamethasone derivative or anti-inflammatory steroid. Furthermore, a wide range of other bioactive materials can be employed, including, but not limited to, the following categories of agents: thrombolytics, vasodilators, antihypertensive agents, antimicrobials or antibiotics, antimitotics, antiproliferatives, antisecretory agents, non-steroidal anti-inflammatory drugs, immunosuppressive agents, growth factors and growth factor antagonists, antitumor and/or chemotherapeutic agents, antipolymerases, antiviral agents, photodynamic therapy agents, antibody targeted therapy agents, prodrugs, sex hormones, free radical scavengers, antioxidants, biologic agents, radiotherapeutic agents, radiopaque agents and radiolabelled agents. The major restriction is that the bioactive material must be able to withstand the coating techniques, for example, the vacuum employed during vapor deposition or plasma deposition of at least one porous layer. In other words, the bioactive material must have a relatively low vapor pressure at the deposition temperature, typically, near or at room temperature.

[0015] The polyamide, parylene or a parylene derivative is applied by catalyst-free vapor deposition.

[0016] "Parylene" is both a generic name for a known group of polymers based on p-xylylene and made by vapor phase polymerization, and a name for the unsubstituted form of the polymer; the latter usage is employed herein. More particularly, parylene or a parylene derivative is created by first heating p-xylylene or a suitable derivative at an appropriate temperature (for example, at about 950°C) to produce the cyclic dimer di-p-xylylene (or a derivative thereof). The resultant solid can be separated in pure form, and then cracked and pyrolyzed at an appropriate temperature (for example, at about 680°C) to produce a monomer vapor of p-xylylene (or derivative); the monomer vapor is cooled to a suitable temperature (for example, below 50°C) and allowed to condense on the desired object, for example, on the at least one layer of bioactive material. The resultant polymer has the repeating structure $\{\text{CH}_2\text{C}_6\text{H}_4\text{CH}_2\}_n$, with n equal to about 5,000, and a molecular weight in the range of 500,000.

[0017] As indicated, parylene and parylene derivative coatings applicable by vapor deposition are known for a variety of biomedical uses, and are commercially available from or through a variety of sources, including Specialty Coating Systems (100 Deposition Drive, Clear Lake, WI 54005), Para Tech Coating, Inc. (35 Argonaut, Aliso Viejo, CA 92656) and Advanced Surface Technology, Inc. (9 Linnet Circle, Billerica, MA 01821-3902).

[0018] The coatings can alternatively be applied by plasma deposition. Plasma is an ionized gas maintained under vacuum and excited by electrical energy, typically in the radiofrequency range. Because the gas is maintained under vacuum, the plasma deposition process occurs at or near room temperature. Plasma can be used to deposit polymers such as poly(ethylene oxide), poly(ethylene glycol), and poly(propylene oxide), as well as polymers of silicone, methane, tetrafluoroethylene (including TEFLON brand polymers), tetramethyldisiloxane, and others.

[0019] While the foregoing represents some preferred embodiments of the present invention, other polymer systems may also be employed, e.g., polymers derived from photopolymerizable monomers. Also, other coating techniques may be utilized, e.g., dipping, spraying, and the like.

[0020] The device may include two or more layers of different bioactive materials atop the structure. However, for the purposes of the present invention, the same bioactive material will generally not be posited on the different surfaces of the device within the same layer. In other words, each surface of the device structure will carry a different bioactive material or materials where the bioactive material is the outermost layer, e.g. heparin may form the innermost layer or the outermost layer or both. These additional layers may be placed directly atop one another or can be separated by additional porous polymer layers between each of them. Additionally, the layers of bioactive materials can comprise a mixture of different bioactive materials. The porous layers are also preferably composed of parylene or a parylene derivative. Advantageously, the two or more bioactive materials can have different solubilities, and the layer containing the less soluble bioactive material (for example, dexamethasone) is preferably posited above the layer containing the more soluble bioactive material (for example, heparin). Unexpectedly, this has been found to increase the *in vitro* release rate of some relatively less soluble materials such as dexamethasone, while simultaneously decreasing the release rate of some relatively more soluble materials such as heparin.

[0021] While the structure included in the device may be configured in a variety of ways, the structure is preferably configured as a vascular stent composed of a biocompatible metal such as stainless steel, nickel, silver, platinum, gold, titanium, tantalum, iridium, tungsten, Nitinol, inconel, or the like. An additional substantially nonporous coating layer of parylene or a parylene derivative or other biocompatible polymer of about 50,000 to 500,000 Å thick may be posited directly atop the vascular stent, beneath the at least one layer of bioactive material. The additional coating layer can merely be relatively less porous than the at least one porous layer, but preferably is substantially nonporous, that is, sufficiently nonporous to render the stent essentially impervious to blood during normal circumstances of use.

[0022] The device and methods of the present invention are useful in a wide variety of locations within a human or veterinary patient, such as in the esophagus, trachea, colon, biliary tract, urinary tract and vascular system, as well as for subdural and orthopedic devices, implants or replacements. They are particularly advantageous for reliably delivering suitable bioactive materials during or following an intravascular procedure, and find particular use in preventing abrupt closure and/or restenosis of a blood vessel. More particularly, they permit, for example, the delivery of an antithrombotic, an antiplatelet, an anti-inflammatory steroid, or another medication to the region of a blood vessel which has been opened by PTA. Likewise, it allows for the delivery of one bioactive material to, for example, the lumen of a blood vessel and another bioactive material to the vessel wall. The use of a porous polymer layer permits the release rate of a bioactive material to be carefully controlled over both the short and long terms.

[0023] These and other aspects of the present invention will be appreciated by those skilled in the art upon the reading and understanding of the specification.

[0024] In another aspect of the invention, the bioactive material is attached to the non-porous layer and is advantageously eluted for prolonged periods of time. The non-porous layer is attached to the base material of the structure. The non-porous layer can be any of those previously or subsequently listed herein, and, likewise, the bioactive material can be any of those previously or subsequently listed herein. Conveniently, and in a preferred embodiment, a glycoprotein IIb/IIIa inhibitor such as commercially available ReoPro™ is attached to a non-porous layer of parylene positioned on the outer surface of the medical device such as a coronary stent. The ReoPro™ is advantageously eluted from the surface of the stent for prolonged periods of time.

Brief Description of the Drawings

[0025] A better understanding of the present invention will now be had upon reference to the following detailed description, when read in conjunction with the accompanying drawing, wherein Fig. 1 to 10 represent prior art known in European Patent application EP 0 747 069 and in which:

FIG. 1 is a cross-sectional view of a first example;

FIG. 2 is a cross-sectional view of another example;

FIG. 3 is a cross-sectional view of yet another example invention;

FIG. 4 is a cross-sectional view of a further example;

FIG. 5 is a cross-sectional view of an additional example;

FIGs. 6A and 6B are cross-sectional views of an additional example;

FIG. 7 is a cross-sectional view of an additional example;

FIG. 8 is a partial, enlarged top view of FIG. 7;

FIG. 9 is an enlarged, sectional view along lines 9-9 of FIG. 8;

FIGs. 10A-10D are enlarged cross-sectional views along lines 10-10 of FIG. 8;

FIG. 11 depicts an aspect of the medical device of the present invention utilizing a polymer coating layer with a bioactive material attached thereto; and

FIG. 12 depicts another aspect of the medical device of FIG. 11 in which the polymer coating layer is adhered to the outer surface of the device base material using an adhesive promotion layer.

Detailed Description

[0026] The example of Figures 1 to 10 are disclosed in the applicant's earlier European Patent application no. EP-A-0,747,069 and are described herein for a better understanding the embodiments covered in the claims of this application in which embodiments are disclosed below with reference to Figures 11 and 12.

[0027] With reference now to FIG. 1, an implantable medical device 10 is shown and first comprises a structure 12 adapted for introduction into a human or veterinary patient. "Adapted" means that the structure 12 is shaped and sized for such introduction. For clarity, only a portion of the structure 12 is shown in FIG. 1.

[0028] By way of example, the structure 12 is configured as a vascular stent particularly adapted for insertion into the vascular system of the patient. However, this stent structure can be used in other systems and sites such as the esophagus, trachea, colon, biliary ducts, urethra and ureters, subdural among others. Indeed, the structure 12 can alternatively be configured as any conventional vascular or other medical device, and can include any of a variety of conventional stents or other adjuncts, such as helical wound strands, perforated cylinders, or the like. Moreover, because the problems addressed by the present invention arise with respect to those portions of the device actually positioned within the patient, the inserted structure 12 need not be an entire device, but can merely be that portion of a vascular or other device which is intended to be introduced into the patient. Accordingly, the structure 12 can be configured as at least one of, or any portion of, a catheter, a wire guide, a cannula, a stent, a vascular or other graft, a cardiac pacemaker lead or lead tip, a cardiac defibrillator lead or lead tip, a heart valve, or an ortho-

pedic device, appliance, implant, or replacement. The structure 12 can also be configured as a combination of portions of any of these.

[0029] Most preferably, however, the structure 12 is configured as a vascular stent such as the commercially available Gianturco-Roubin FLEX-STENT® or GR II™ coronary stent from Cook Incorporated, Bloomington, Indiana. Such stents are typically about 10 to about 60 mm in length and designed to expand to a diameter of about 2 to about 6 mm when inserted into the vascular system of the patient. The Gianturco-Roubin stent in particular is typically about 12 to about 25 mm in length and designed to expand to a diameter of about 2 to about 4 mm when so inserted.

[0030] These stent dimensions are, of course, applicable to exemplary stents employed in the coronary arteries. Structures such as stents or catheter portions intended to be employed at other sites in the patient, such as in the aorta, esophagus, trachea, colon, biliary tract, or urinary tract will have different dimensions more suited to such use. For example, aortic, esophageal, tracheal and colonic stents may have diameters up to about 25 mm and lengths about 100 mm or longer.

[0031] The structure 12 is composed of a base material 14 suitable for the intended use of the structure 12. The base material 14 is preferably biocompatible, although cytotoxic or other poisonous base materials may be employed if they are adequately isolated from the patient. Such incompatible materials may be useful in, for example, radiation treatments in which a radioactive material is positioned by catheter in or close to the specific tissues to be treated. Under most circumstances, however, the base material 14 of the structure 12 should be biocompatible.

[0032] A variety of conventional materials can be employed as the base material 14. Some materials may be more useful for structures other than the coronary stent exemplifying the structure 12. The base material 14 may be either elastic or inelastic, depending upon the flexibility or elasticity of the polymer layers to be applied over it. The base material may be either biodegradable or non-biodegradable, and a variety of biodegradable polymers are known. Moreover, some biologic agents have sufficient strength to serve as the base material 14 of some useful structures 12, even if not especially useful in the exemplary coronary stent.

[0033] Accordingly, the base material 14 can include at least one of stainless steel, tantalum, titanium, nitinol, gold, platinum, inconel, iridium, silver, tungsten, or another biocompatible metal, or alloys of any of these; carbon or carbon fiber; cellulose acetate, cellulose nitrate, silicone, polyethylene terephthalate, polyurethane, polyamide, polyester, polyorthoester, polyanhydride, polyether sulfone, polycarbonate, polypropylene, high molecular weight polyethylene, polytetrafluoroethylene, or another biocompatible polymeric material, or mixtures or copolymers of these; polylactic acid, polyglycolic acid or copolymers thereof, a polyanhydride, polyc-

aprolactone, polyhydroxybutyrate valerate or another biodegradable polymer, or mixtures or copolymers of these; a protein, an extracellular matrix component, collagen, fibrin or another biologic agent; or a suitable mixture of any of these. Stainless steel is particularly useful as the base material 14 when the structure 12 is configured as a vascular stent.

[0034] Of course, when the structure 12 is composed of a radiolucent material such as polypropylene, polyethylene, or others above, a conventional radiopaque coating may and preferably should be applied to it. The radiopaque coating provides a means for identifying the location of the structure 12 by X-ray or fluoroscopy during or after its introduction into the patient's vascular system.

[0035] With continued reference to FIG. 1, the vascular device 10 next comprises at least one layer 18 of a bioactive material posited on one surface of the structure 12. For the purposes of the present invention, at least one bioactive material is posited on one surface of the structure 12, and the other surface will either contain no bioactive material or one or more different bioactive materials. In this manner, one or more bioactive materials or drugs may be delivered, for example, with a vascular stent, to the blood stream from the lumen surface of the stent, and a different treatment may be delivered on the vessel surface of the stent. A vast range of drugs, medicaments and materials may be employed as the bioactive material in the layer 18, so long as the selected material can survive exposure to the vacuum drawn during vapor deposition or plasma deposition. Particularly useful in the practice of the present invention are materials which prevent or ameliorate abrupt closure and restenosis of blood vessels previously opened by stenting surgery or other procedures. Thrombolytics (which dissolve, break up or disperse thrombi) and antithrombotics (which interfere with or prevent the formation of thrombi) are especially useful bioactive materials when the structure 12 is a vascular stent. Particularly preferred thrombolytics are urokinase, streptokinase, and the tissue plasminogen activators. Particularly preferred antithrombotics are heparin, hirudin, and the antiplatelets.

[0036] Urokinase is a plasminogen activating enzyme typically obtained from human kidney cell cultures. Urokinase catalyzes the conversion of plasminogen into the fibrinolytic plasmin, which breaks down fibrin thrombi.

[0037] Heparin is a mucopolysaccharide anticoagulant typically obtained from porcine intestinal mucosa or bovine lung. Heparin acts as a thrombin inhibitor by greatly enhancing the effects of the blood's endogenous antithrombin III. Thrombin, a potent enzyme in the coagulation cascade, is key in catalyzing the formation of fibrin. Therefore, by inhibiting thrombin, heparin inhibits the formation of fibrin thrombi. Alternatively, heparin may be covalently bound to the outer layer of structure 12. Thus, heparin would form the outermost layer of

structure 12 and would not be readily degraded enzymatically, and would remain active as a thrombin inhibitor.

[0038] Of course, bioactive materials having other functions can also be successfully delivered by the device 10. For example, an antiproliferative agent such as methotrexate will inhibit over-proliferation of smooth muscle cells and thus inhibit restenosis of the dilated segment of the blood vessel. The antiproliferative is desirably supplied for this purpose over a period of about four to six months. Additionally, localized delivery of an antiproliferative agent is also useful for the treatment of a variety of malignant conditions characterized by highly vascular growth. In such cases, the device 10 could be placed in the arterial supply of the tumor to provide a means of delivering a relatively high dose of the antiproliferative agent directly to the tumor.

[0039] A vasodilator such as a calcium channel blocker or a nitrate will suppress vasospasm, which is common following angioplasty procedures. Vasospasm occurs as a response to injury of a blood vessel, and the tendency toward vasospasm decreases as the vessel heals. Accordingly, the vasodilator is desirably supplied over a period of about two to three weeks. Of course, trauma from angioplasty is not the only vessel injury which can cause vasospasm, and the device 10 may be introduced into vessels other than the coronary arteries, such as the aorta, carotid arteries, renal arteries, iliac arteries or peripheral arteries for the prevention of vasospasm in them.

[0040] A variety of other bioactive materials are particularly suitable for use when the structure 12 is configured as something other than a coronary stent. For example, an anti-cancer chemotherapeutic agent can be delivered by the device 10 to a localized tumor. More particularly, the device 10 can be placed in an artery supplying blood to the tumor or elsewhere to deliver a relatively high and prolonged dose of the agent directly to the tumor, while limiting systemic exposure and toxicity. The agent may be a curative, a pre-operative debulker reducing the size of the tumor, or a palliative which eases the symptoms of the disease. It should be noted that the bioactive material in the present invention is delivered across the device 10, and not by passage from an outside source through any lumen defined in the device 10, such as through a catheter employed for conventional chemotherapy. The bioactive material of the present invention may, of course, be released from the device 10 into any lumen defined in the device, or to tissue in contact with the device and that the lumen may carry some other agent to be delivered through it. For example, tamoxifen citrate, Taxol® or derivatives thereof Proscar®, Hytrin®, or Eulexin® may be applied to the tissue-exposed surface of the device for delivery to a tumor located, for example in breast tissue or the prostate.

[0041] Dopamine or a dopamine agonist such as bromocriptine mesylate or pergolide mesylate is useful for

the treatment of neurological disorders such as Parkinson's disease. The device 10 could be placed in the vascular supply of the thalamic substantia nigra for this purpose, or elsewhere, localizing treatment in the thalamus.

[0042] A wide range of other bioactive materials can be delivered by the device 10. Accordingly, it is preferred that the bioactive material contained in the layer 18 includes at least one of heparin, covalent heparin, or another thrombin inhibitor, hirudin, hirulog, argatroban, D-phenylalanyl-L-poly-L-arginyl chloromethyl ketone, or another antithrombogenic agent, or mixtures thereof; urokinase, streptokinase, a tissue plasminogen activator, or another thrombolytic agent, or mixtures thereof; a fibrinolytic agent; a vasospasm inhibitor; a calcium channel blocker, a nitrate, nitric oxide, a nitric oxide promoter or another vasodilator; Hytrin® or other antihypertensive agents; an antimicrobial agent or antibiotic; aspirin, ticlopidine, a glycoprotein IIb/IIIa inhibitor or another inhibitor of surface glycoprotein receptors, or another antiplatelet agent; colchicine or another antimitotic, or another microtubule inhibitor, dimethyl sulfoxide (DMSO), a retinoid or another antisecretory agent; cytochalasin or another actin inhibitor; or a remodelling inhibitor; deoxyribonucleic acid, an antisense nucleotide or another agent for molecular genetic intervention; methotrexate or another antimetabolite or antiproliferative agent; tamoxifen citrate, Taxol® or the derivatives thereof, or other anti-cancer chemotherapeutic agents; dexamethasone, dexamethasone sodium phosphate, dexamethasone acetate or another dexamethasone derivative, or another anti-inflammatory steroid or non-steroidal antiinflammatory agent; cyclosporin or another immunosuppressive agent; trapidil (a PDGF antagonist), angiopeptin (a growth hormone antagonist), angiotensin, a growth factor or an anti-growth factor antibody, or another growth factor antagonist; dopamine, bromocriptine mesylate, pergolide mesylate or another dopamine agonist; ⁶⁰Co (5.3 year half life), ¹⁹²Ir (73.8 days), ³²P (14.3 days), ¹¹¹In (68 hours), ⁹⁰Y (64 hours), ^{99m}Tc (6 hours) or another radiotherapeutic agent; iodine-containing compounds, barium-containing compounds, gold, tantalum, platinum, tungsten or another heavy metal functioning as a radiopaque agent; a peptide, a protein, an enzyme, an extracellular matrix component, a cellular component or another biologic agent; captopril, enalapril or another angiotensin converting enzyme (ACE) inhibitor; ascorbic acid, alpha tocopherol, superoxide dismutase, deferoxamine, a 21-aminosteroid (lasaroid) or another free radical scavenger, iron chelator or antioxidant; a ¹⁴C-, ³H-, ¹³¹I-, ³²P- or ³⁶S-radiolabelled form or other radiolabelled form of any of the foregoing; estrogen or another sex hormone; AZT or other antipolymerases; acyclovir, famciclovir, rimantadine hydrochloride, ganciclovir sodium, Norvir, Crixivan, or other antiviral agents; 5-aminolevulinic acid, meta-tetrahydroxyphenylchlorin, hexadecafluoro zinc phthalocyanine, tetramethyl he-

matoporphyrin, rhodamine 123 or other photodynamic therapy agents; an IgG2 Kappa antibody against *Pseudomonas aeruginosa* exotoxin A and reactive with A431 epidermoid carcinoma cells, monoclonal antibody against the noradrenergic enzyme dopamine beta-hydroxylase conjugated to saporin or other antibody targeted therapy agents; gene therapy agents; and enalapril and other prodrugs; Proscar®, Hytrin® or other agents for treating benign prostatic hyperplasia (BHP) or a mixture of any of these; and various forms of small intestine submucosa (SIS).

[0043] In a particularly preferred aspect, the layer of bioactive material contains preferably from about 0.01 mg to about 10 mg and more preferably from about 0.1 mg to about 4 mg of the bioactive material per cm² of the gross surface area of the structure. "Gross surface area" refers to the area calculated from the gross or overall extent of the structure, and not necessarily to the actual surface area of the particular shape or individual parts of the structure. In other terms, about 100 µg to about 300 µg of drug per 0.001 inch of coating thickness may be contained on the device surface.

[0044] When the structure 12 is configured as a vascular stent, however, particularly preferred materials for the bioactive material of the layer 18 are heparin, anti-inflammatory steroids including but not limited to dexamethasone and its derivatives, and mixtures of heparin and such steroids.

[0045] Still with reference to FIG. 1, the device 10 also comprises at least one porous layer 20 posited over the layer 18 of bioactive material and the bioactive-material-free surface. The purpose of the porous layer 20 is to provide a controlled release of the bioactive material when the device 10 is positioned in the vascular system of a patient. The thickness of the porous layer 20 is chosen so as to provide such control.

[0046] More particularly, the porous layer 20 is composed of a polymer deposited on the bioactive material layer 18, preferably by vapor deposition. Plasma deposition may also be useful for this purpose. Preferably, the layer 20 is one that is polymerized from a vapor which is free of any solvent, catalysts or similar polymerization promoters. Also preferably, the polymer in the porous layer 20 is one which automatically polymerizes upon condensation from the vapor phase, without the action of any curative agent or activity such as heating, the application of visible or ultraviolet light, radiation, ultrasound, or the like. Most preferably, the polymer in the porous layer 20 is polyimide, parylene or a parylene derivative.

[0047] When first deposited, the parylene or parylene derivative is thought to form a network resembling a fibrous mesh, with relatively large pores. As more is deposited, the porous layer 20 not only becomes thicker, but it is believed that parylene or parylene derivative is also deposited in the previously formed pores, making the existing pores smaller. Careful and precise control over the deposition of the parylene or parylene deriva-

tive therefore permits close control over the release rate of material from the at least one layer 18 of bioactive material. It is for this reason that the bioactive material lies under the at least one porous layer 20, rather than being dispersed within or throughout it. The porous layer 20, however, also protects the bioactive material layer 18 during deployment of the device 10, for example, during insertion of the device 10 through a catheter and into the vascular system or elsewhere in the patient.

[0048] As shown in FIG. 1, the device 10 can further comprise at least one additional coating layer 16 posited between the structure 12 and the at least one layer 18 of bioactive material. While the additional coating layer 16 can simply be a medical grade primer, the additional coating layer 16 is preferably composed of the same polymer as the at least one porous layer 20. However, the additional coating layer 16 is also preferably less porous than the at least one porous layer 20, and is more preferably substantially nonporous. "Substantially nonporous" means that the additional coating layer 16 is sufficiently impervious to prevent any appreciable interaction between the base material 14 of the structure 12 and the blood to which the device 10 will be exposed during use. The use of an additional coating layer 16 which is substantially nonporous would permit the use of a toxic or poisonous base material 14, as mentioned above. Even if the base material 14 of the structure 12 is biocompatible, however, it may be advantageous to isolate it from the blood by use of a substantially nonporous coating layer 16.

[0049] Other polymer systems that may find application within the scope of the invention include polymers derived from photopolymerizable monomers such as liquid monomers preferably having at least two cross linkable C-C (Carbon to Carbon) double bonds and being a non-gaseous addition polymerizable ethylenically unsaturated compound, having a boiling point above 100°C, at atmospheric pressure, a molecular weight of about 100-1500 and being capable of forming high molecular weight addition polymers readily. More preferably, the monomer is preferably an addition photopolymerizable polyethylenically unsaturated acrylic or methacrylic acid ester containing two or more acrylate or methacrylate groups per molecule or mixtures thereof. A few illustrative examples of such multifunctional acrylates are ethylene glycol diacrylate, ethylene glycol dimethacrylate, trimethylpropane triacrylate, trimethylpropane trimethacrylate, pentaerythritol tetraacrylate or pentaerythritol tetramethacrylate, 1,6-hexanediol dimethacrylate, and diethyleneglycol dimethacrylate.

[0050] Also useful in some special instances are monoacrylates such as n-butylacrylate, n-butyl methacrylate, 2-ethylhexyl acrylate, lauryl-acrylate, and 2-hydroxypropyl acrylate. Small quantities of amides of (meth)acrylic acid such as N-methylol methacrylamide butyl ether are also suitable; N-vinyl compounds such as N-vinyl pyrrolidone, vinyl esters of aliphatic monocarboxylic acids such as vinyl oleate, vinyl ethers of diols

such as butanediol-1, 4-divinyl ether and allyl ether and allyl ester are also suitable. Also included would be other monomers such as the reaction products of di- or polyepoxides such as butanediol-1, 4-diglycidyl ether or bisphenol A diglycidyl ether with (meth)acrylic acid. The characteristics of the photopolymerizable liquid dispersing medium can be modified for the specific purpose by a suitable selection of monomers or mixtures thereof.

[0051] Other useful polymer systems include a polymer that is biocompatible and minimizes irritation to the vessel wall when the stent is implanted. The polymer may be either a biostable or a bioabsorbable polymer depending on the desired rate of release or the desired degree of polymer stability, but a bioabsorbable polymer is preferred for this embodiment since, unlike a biostable polymer, it will not be present long after implantation to cause any adverse, chronic local response. Bioabsorbable polymers that could be used include poly(L-lactic acid), polycaprolactone, poly(lactide-co-glycolide), poly(hydroxybutyrate), poly(hydroxybutyrate-co-valerate), polydioxanone, polyorthoester, polyanhydride, poly(glycolic acid), poly(D,L-lactic acid), poly(glycolic acid-co-trimethylene carbonate), polyphosphoester, polyphosphoester urethane, poly(amino acids), cyanoacrylates, poly(trimethylene carbonate), poly(iminocarbonate), copoly(ether-esters) (e.g., PEO/PLA), polyalkylene oxalates, polyphosphazenes and biomolecules such as fibrin, fibrinogen, cellulose, starch, collagen and hyaluronic acid. Also, biostable polymers with a relatively low chronic tissue response such as polyurethanes, silicones, and polyesters could be used and other polymers could also be used if they can be dissolved and cured or polymerized on the stent such as polyolefins, polyisobutylene and ethylene-alphaolefin copolymers; acrylic polymers and copolymers, vinyl halide polymers and copolymers, such as polyvinyl chloride; polyvinyl ethers, such as polyvinyl methyl ether; polyvinylidene halides, such as polyvinylidene fluoride and polyvinylidene chloride; polyacrylonitrile, polyvinyl ketones; polyvinyl aromatics, such as polystyrene, polyvinyl esters, such as polyvinyl acetate; copolymers of vinyl monomers with each other and olefins, such as ethylene-methyl methacrylate copolymers, acrylonitrile-styrene copolymers, ABS resins, and ethylene-vinyl acetate copolymers; polyamides, such as Nylon 66 and polycaprolactam; alkyd resins, polycarbonates; polyoxymethylenes; polyimides; polyethers; epoxy resins, polyurethanes; rayon; rayon-triacetate; cellulose, cellulose acetate, cellulose butyrate; cellulose acetate butyrate; cellophane; cellulose nitrate; cellulose propionate; cellulose ethers; and carboxymethyl cellulose.

[0052] While plasma deposition and vapor phase deposition may be a preferred method for applying the various coatings on the stent surfaces, other techniques may be employed. For example, a polymer solution may be applied to the stent and the solvent allowed to evaporate, thereby leaving on the stent surface a coating of the polymer and the therapeutic substance. Typically,

the solution can be applied to the stent by either spraying the solution onto the stent or immersing the stent in the solution. Whether one chooses application by immersion or application by spraying depends principally on the viscosity and surface tension of the solution, however, it has been found that spraying in a fine spray such as that available from an airbrush will provide a coating with the greatest uniformity and will provide the greatest control over the amount of coating material to be applied to the stent. In either a coating applied by spraying or by immersion, multiple application steps are generally desirable to provide improved coating uniformity and improved control over the amount of therapeutic substance to be applied to the stent.

[0053] When the layer 18 of bioactive material contains a relatively soluble material such as heparin, and when the at least one porous layer 20 is composed of parylene or a parylene derivative, the at least one porous layer 20 is preferably about 5,000 to 250,000 Å thick, more preferably about 5,000 to 100,000 Å thick, and optimally about 50,000 Å thick. When the at least one additional coating layer 16 is composed of parylene or a parylene derivative, the at least one additional coating is preferably about 50,000 to 500,000 Å thick, more preferably about 100,000 to 500,000 Å thick, and optimally about 200,000 Å thick.

[0054] When the at least one layer 18 of bioactive material contains a relatively soluble material such as heparin, the at least one layer 18 preferably contains a total of about 0.1 to 4 mg of bioactive material per cm² of the gross surface area of the structure 12. This provides a release rate for the heparin (measured in vitro) which is desirably in the range of 0.1 to 0.5 mg/cm² per day, and preferably about 0.25 mg/cm² per day, under typical blood flows through vascular stents. It should be noted that the solubility of dexamethasone can be adjusted as desired, with or without the inclusion of heparin, by mixing it with one or more of its relatively more soluble derivatives, such as dexamethasone sodium phosphate.

[0055] FIG. 11 depicts an aspect of device 10 of the present invention in which coating layer 16 is applied directly to the outer surface of base material 14 of structure 12. In this configuration, coating layer 16 is preferably a non-porous coating layer as previously described. When coating layer 16 comprises a parylene derivative, non-porous coating layer 16 ranges in thickness from 50,000 to 500,000 Å, more preferably in the range of 100,000 to 500,000 Å, and optimally about 200,000 Å. In this aspect of the invention, non-porous coating layer 16 is also an adsorbent in which an adsorbent is defined as an agent that attracts other materials or particles to its surface as indicated in Dorland's Illustrated Medical Dictionary 26th Edition by W.B. Saunders Co., Philadelphia, PA. Bioactive material 18 is then attached to the surface of coating layer 16.

[0056] Alternatively, alternating layers of coating material and the same or different bioactive materials can

be applied to the surface of bioactive material layer 18. However, in this particular aspect of the invention, the outer layer of structure 12 is a bioactive material layer 18.

[0057] In another aspect of the present invention as depicted in FIG. 11, coating layer 16 can be considered an adsorbent layer and/or an absorbent layer in which a bioactive material is attached thereto. In one particular example, device 10 is a stainless steel GR II™ stent in which the stainless steel base material 14 of structure 12 is coated with a polymer and, in particular, parylene. This adsorbent polymer layer 16 of parylene is approximately 230,000 Å thick. Bioactive material layer 18 of the antiplatelet GP IIb/IIIa antibody (AZ1) was passively loaded on adsorbent polymer layer 16. The polymer coated stainless steel stents were immersed for approximately 24 hours in a buffered aqueous solution of AZ1 antibody (1 mg/ml, pH = 7.2) at 37°C. AZ1 is a monoclonal anti-rabbit platelet glycoprotein (GP) IIb/IIIa antibody. Using radio-labeled AZ1, it was demonstrated that approximately 0.02 µg antibody was loaded per mm² stent surface area (approximately 2 µg total for a 3 x 20 mm GR II™ stent). It was also demonstrated that in an in-vitro flow system (10 ml/min, 1% BSA in PBS) approximately half the loaded antibody remained on the stent after approximately 10 days perfusion.

[0058] The mechanism by which the stent is loaded with drug are thought to include adsorption onto the surface of the polymer layer and/or absorption into the polymer.

[0059] Previous studies with similar loading and release of AZ1 from cellulose coated stainless steel stents showed inhibition of platelet aggregation and reduced thrombosis rates in a rabbit model of deep arterial injury. (Aggarwal, et al., Antithrombotic Potential of Polymer-Coated Stents Eluting Platelet Glycoprotein IIb/IIIa Receptor Antibody, American Heart Association Circulation Vol. 94 No. 12, December 15, 1996, pp3311-3317).

[0060] In another example, c7E3 Fab as bioactive material layer 18 is attached to polymer coating layer 16. Bioactive material c7E3 Fab is a chimeric monoclonal antibody that acts upon the Gp IIa/IIIb integrin on platelets to inhibit their aggregation. This antibody or receptor blocker can be used in humans intravenously to prevent thrombosis during coronary angioplasty. This receptor blocker is also known as ReoPro™ available from Eli Lilly, Indianapolis, IN. Bioactive material layer 18 of the antiplatelet GP IIb/IIIa antibody (c7E3 Fab) was passively loaded on adsorbent polymer layer 16. The polymer coated stainless steel stents were immersed for approximately 24 hours in a buffered aqueous solution of c7E3 Fab antibody (1 mg/ml, pH = 7.2) at 37°C. c7E3 Fab is an inhibitor of platelet thrombus in humans. Using radio-labeled c7E3 Fab, it was demonstrated that approximately 0.010 µg antibody was loaded per mm² stent surface area (approximately 10 µg total for a 3 x 20 mm GR II™ stent). It was also demonstrated that in

an in-vitro flow system (10 ml/min, 1% BSA in PBS) approximately half the loaded antibody remained on the stent after approximately 10 days perfusion.

[0061] FIG. 12 depicts still another aspect of device 10 of FIG. 11. In this embodiment, a parylene adhesion promotion layer 30 is first applied to stainless steel base material 14 of structure 12. By way of example, adhesion promotion layer 30 is a thin layer of silane having a thickness in the range of, for example, 0.5 to 5,000 Å and preferably, 2 to 50 Å. This silane promotion layer is preferably A-174 silane including a gamma-methacryloxypropyltrimethoxysilane, which is available from Specialty Coating Systems Inc., Indianapolis, IN. In preparing the outer surface of base material 14, it is first cleaned with isopropyl alcohol. The stent is then dipped in the silane to apply a very thin layer thereof to the outer surface of the base material. Polymer coating layer 16 of parylene is then applied to the silane layer. Other methods of preparing the outer surface of base material 14 include plasma etching and grit blasting. Preparation includes cleaning the outer surface of the base material with isopropyl alcohol, plasma etching the outer surface of the base material and applying the silane to the plasma etched surface. With grit blasting, the outer surface of the base material is grit blasted and then cleaned with isopropyl alcohol to which silane is applied to the cleansed grit blasted surface.

[0062] As shown in FIG. 2, the device 10 is not limited to the inclusion of a single layer 18 of bioactive material. The device 10 can, for example, comprise a second layer 22 of a bioactive material posited over the structure 12. The bioactive material of the second layer 22 can be, but need not necessarily be, different from the bioactive material of the first bioactive material layer 18, only that they not be posited on the same surface of the device 10 without the intermediate porous layer 24. The use of different materials in the layers 18 and 22 allows the device 10 to perform more than a single therapeutic function.

[0063] The device 10 can further comprise an additional porous layer 24 of the polymer posited between each of the layers 18 and 22 of bioactive material. It is reiterated that bioactive material 18 is on one surface of structure 12. The other surface may be free of bioactive material or may comprise one or more different bioactive materials. The additional porous layer 24 can give the bioactive materials in the layers 18 and 22 different release rates. Simultaneously, or alternatively, the device 10 may employ bioactive materials in the two layers 18 and 22 which are different from one another and have different solubilities. In such a case, it is advantageous and preferred to position the layer 22 containing the less soluble bioactive material above the layer 18 containing the more soluble bioactive material. Alternatively, the bioactive material 18 may be contained in holes, wells, slots and the like occurring within the stent surface as illustrated in FIGs. 8-10 and will further be discussed in greater detail.

[0064] The device 10 need not include the additional porous layer 24 when two or more layers 18 and 22 of bioactive material are present. As shown in FIG. 3, the layers 18 and 22 do not have to be separated by a porous layer, but can instead lie directly against one another. It is still advantageous in this embodiment to position the layer 22 containing the relatively less soluble bioactive material above the layer 18 containing the relatively more soluble bioactive material.

[0065] Whether or not the additional porous layer 24 is present, it is preferred that the layers 18 and 22 contain about 0.05 to 2.0 mg of each of heparin and dexamethasone, respectively, per 1 cm² of the gross surface area of the structure 12. The total amount of bioactive material posited in the layers 18 and 22 over the structure 12 is thus preferably in the range of about 0.1 to 10 mg/cm².

[0066] The particular structure of the device 10 as disclosed may be adapted to specific uses in a variety of ways. For example, the device 10 may include further layers of the same or different bioactive materials. These additional layers of bioactive material may or may not be separated by additional porous layers, as convenient or desired. Alternatively, additional porous layers may separate only some of the additional layers of bioactive material. Moreover, one bioactive material may be placed on one portion of the structure 12 of the device 10, and another bioactive material placed on a different portion of the structure 12 of the device 10.

[0067] Alternatively, the device 10 need not include the additional coating layer 16 at all. Such a configuration is shown in FIG. 4, in which the bioactive material layer 18 is posited directly atop the base material 14 of the structure 12. In such a case, it may be highly advantageous to surface process or surface activate the base material 14, to promote the deposition or adhesion of the bioactive material on the base material 14. Surface processing and surface activation can also selectively alter the release rate of the bioactive material. Such processing can also be used to promote the deposition or adhesion of the additional coating layer 16, if present, on the base material 14. The additional coating layer 16 itself, or any second or additional porous layer 24 itself, can similarly be processed to promote the deposition or adhesion of the bioactive material layer 18, or to further control the release rate of the bioactive material.

[0068] Useful methods of surface processing can include any of a variety of such procedures, including: cleaning; physical modifications such as etching, drilling, cutting, or abrasion; and chemical modifications such as solvent treatment, the application of primer coatings, the application of surfactants, plasma treatment, ion bombardment and covalent bonding.

[0069] It has been found particularly advantageous to plasma treat the additional coating layer 16 (for example, of parylene) before depositing the bioactive material layer 18 atop it. The plasma treatment improves the adhesion of the bioactive material, increases the amount

of bioactive material that can be deposited, and allows the bioactive material to be deposited in a more uniform layer. Indeed, it is very difficult to deposit a hygroscopic agent such as heparin on an unmodified parylene surface, which is hydrophobic and poorly wettable. However, plasma treatment renders the parylene surface wettable, allowing heparin to be easily deposited on it.

[0070] Referring now to FIG. 5, an example of the device 10 is there shown in which a mechanical bond or connector 26 is provided between (a) any one of the porous layers 20 and 24, and (b) any or all of the other of the porous layers 20 and 24, the additional coating layer 16 and the base material 14. The connector 26 reliably secures the layers 16, 20 and/or 24 to each other, and or to the base material 14. The connector 26 lends structural integrity to the device 10, particularly after the bioactive material layer or layers 18 and/or 20 have been fully released into the patient.

[0071] For simplicity, the connector 26 is shown in FIG. 5 as a plurality of projections of the base material 14 securing a single porous layer 20 to the base material 14. The connector 26 may alternatively extend from the porous layer 20, through the bioactive material layer 18, and to the base material 14. In either case, a single layer 18 of bioactive material, divided into several segments by the connector 26, is posited between the porous layer 20 and the base material 14. The connectors can also function to partition the different bioactive agents into different regions of the device's surface.

[0072] The connector 26 may be provided in a variety of ways. For example, the connector 26 can be formed as a single piece with the base material 14 during its initial fabrication or molding into the structure 12. The connector 26 can instead be formed as a distinct element, such as a bridge, strut, pin or stud added to an existing structure 12. The connector 26 can also be formed as a built-up land, shoulder, plateau, pod or pan on the base material 14. Alternatively, a portion of the base material 14 between the desired locations of plural connectors 26 may be removed by etching, mechanical abrasion, or the like, and the bioactive material layer 18 deposited between them. The connector 26 can also be formed so as to extend downwards towards the base material 14, by wiping or etching away a portion of a previously applied bioactive material layer 18, and allowing the porous layer 20 to deposit by vapor deposition or plasma deposition directly on the bare portions of the base material 14. Other ways to expose a portion of the base material 14 to direct connection to the porous layer 20 will be evident to those skilled in this area.

[0073] In another example, as illustrated in FIGs. 6A, 6B and 7, a bioactive material 18 is posited on the one surface of base material 14 making up structure 12 in FIG. 6A. FIG. 7 shows a stent 10 in its flat or planar state prior to being coiled and showing porous layer 20 applied to its outermost surface. FIGs. 6A and 6B are section views along line 6-6 of FIG. 7. The bioactive material 18 posited on the one surface of base material 14 in FIG.

6A may be a number of different therapeutic and/or diagnostic agents. For example, the device 10 may be a stent which is placed in the body of a patient near a tumor to deliver a chemotherapeutic agent, such as tamoxifen citrate or Taxol®, directly to the tumor. A porous layer 20 is posited over the bioactive material 18 to provide a smoother surface as well as a more controlled release of the bioactive material 18. As further illustrated in FIG. 6A, the opposite surface of the device 10 may have, for example, heparin 18' covalently bonded to porous layer 20, particularly where this surface faces, for example, the lumen of a blood vessel, to provide antithrombotic effect and blood compatibility. It is pointed out, as has been discussed herein, a third but different bioactive material may be posited (not shown) on the opposite surface of base material 14 from the first bioactive material 18 and on the same side of base material 14 as the covalently bound heparin or any other bioactive material including other covalently bound bioactive materials and separated by porous layer 20.

[0074] A variation of the example shown in FIG. 6A is illustrated in FIG. 6B, where two bioactive materials 18 and 18' are posited on the same surface of base material 14 of structure 12. A porous layer 20 may be deposited over the bioactive materials 18 and 18' as well as the bioactive-material-free surface of base material 14. This example illustrates a situation where it may be desirable to deliver two agents to the tissue to which the particular surface of device 10 is exposed, e.g., an anti-inflammatory agent and an antiviral agent. Moreover, the opposite surface of the device free of bioactive material is available for positing one or more bioactive materials or therapeutic agents, e.g., an antithrombotic agent.

[0075] As has been previously discussed, multiple layers of bioactive materials and porous layers may be applied to the device 10 where the limiting factors become the total thickness of the device, the adhesion of multiple layers and the like.

[0076] In still another example, the device includes apertures within the device for containing the bioactive material. This example is illustrated in FIGs. 8, 9, 10A, 10B, 10C and 10D. FIG. 8 shows an arm of the stent of FIG. 7 wherein the arm includes holes 28 into which a bioactive material is contained. FIG. 9 shows a section of the arm of the stent along lines 9-9 of FIG. 8. Bioactive material 18 is contained within the hole 28 where the base material 14 contains coating 16 and further where porous layer 20 forms the outer most layer for the bioactive material 18 to diffuse through. In an alternative embodiment, wells 28' may be cut, etched or stamped into the base material 14 of the device in which a bioactive material 18 may be contained. This embodiment is illustrated in FIGs. 10A, 10B, 10C and 10D which are sectional FIGs. taken along line 10-10 of FIG. 8. The wells 28' may also be in the form of slots or grooves in the surface of the base material 14 of the medical device. This example provides the advantage of better

controlling the total amount of the bioactive material 18 to be released as well as the rate at which it is released. For example, a V-shape well 28', as illustrated in FIG. 10D, will contain less quantity of bioactive material 18 and release the material at geometric rate as compared to a square shaped well 28', as illustrated in FIG. 10B, which will have a more uniform, linear release rate.

[0077] The holes, wells, slots, grooves and the like, described above, may be formed in the surface of the device 10 by a variety of techniques. For example, such techniques include drilling or cutting by utilizing lasers, electron-beam machining and the like or employing photoresist procedures and etching the desired apertures.

[0078] All the bioactive materials discussed above that may be coated on the surface of the device 10 may be used to be contained within the apertures. Likewise, layers of bioactive materials and porous layers may be applied and built up on the exterior surfaces of the device as described previously, e.g., heparin, may be covalently bound to one surface of the device illustrated in FIG. 9.

[0079] The method of making the device 10 according to the example of figures 1 to 10 may now be understood. In its simplest form, the method comprises the steps of depositing the at least one layer 18 of bioactive material over the structure 12, followed by depositing the at least one porous layer 20, preferably by vapor deposition or plasma deposition, over the at least one bioactive material layer 18 on the one surface of structure 12. The at least one porous layer 20 being composed of a biocompatible polymer and being of a thickness adequate to provide a controlled release of the bioactive material. Preferably, the at least one additional coating layer 16 is first posited by vapor deposition directly on the base material 14 of the structure 12. Such deposition is carried out by preparing or obtaining di-p-xylylene or a derivative thereof, sublimating and cracking the di-p-xylylene or derivative to yield monomeric p-xylylene or a monomeric derivative, and allowing the monomer to simultaneously condense on and polymerize over the base material 14. The deposition step is carried out under vacuum, and the base material 14 maintained at or near room temperature during the deposition step. The deposition is carried out in the absence of any solvent or catalyst for the polymer, and in the absence of any other action to aid polymerization. One preferred derivative for carrying out the deposition step is dichloro-di-p-xylylene. The parylene or parylene derivative is preferably applied at the thickness disclosed above, to yield a coating layer 16 which is substantially nonporous, but in any event less porous than the at least one porous layer 20 to be applied. If required by the composition of the coating layer 16, the layer 16 is then surface processed in an appropriate manner, for example, by plasma treatment as disclosed above.

[0080] The at least one layer 18 of the desired bioactive material or materials is then applied to the one surface of the structure 12, and in particular, onto the ad-

ditional coating layer 16. This application step can be carried out in any of a variety of convenient ways, such as by dipping, rolling, brushing or spraying a fluid mixture of the bioactive material onto the additional coating layer 16, or by electrostatic deposition of either a fluid mixture or dry powder of the bioactive material, or by any other appropriate method. Different bioactive agents may be applied to different sections or surfaces of the device:

[0081] It can be particularly convenient to apply a mixture of the bioactive material or materials and a volatile fluid over the structure, and then remove the fluid in any suitable way, for example, by allowing it to evaporate. When heparin and/or dexamethasone or its derivatives serve as the bioactive material(s), the fluid is preferably ethyl alcohol. The bioactive material is preferably applied in an amount as disclosed above.

[0082] Other methods of depositing the bioactive material layer 18 over the structure 12 would be equally useful.

[0083] The bioactive material may, of course, be deposited on the one surface of the structure 12 as a smooth film or as a layer of particles. Moreover, multiple but different bioactive materials may be deposited in a manner that different surfaces of the device contain the different bioactive agents. In the latter case, the particle size may affect the properties or characteristics of the device 10, such as the smoothness of the uppermost porous coating 20, the profile of the device 10, the surface area over which the bioactive material layer 18 is disposed, the release rate of the bioactive material, the formation of bumps or irregularities in the bioactive material layer 18, the uniformity and strength of adhesion of the bioactive material layer 18, and other properties or characteristics. For example, it has been useful to employ micronized bioactive materials, that is, materials which have been processed to a small particle size, typically less than 10µm in diameter. However, the bioactive material may also be deposited as microencapsulated particles, dispersed in liposomes, adsorbed onto or absorbed into small carrier particles, or the like.

[0084] In still another embodiment, the bioactive material may be posited on the one surface of structure 12 in a specific geometric pattern. For example, the tips or arms of a stent may be free of bioactive material, or the bioactive material may be applied in parallel lines, particularly where two or more bioactive materials are applied to the same surface.

[0085] Any other layers, such as the second bioactive material layer 22 or the additional porous layer 24, are applied in the appropriate order and in the same manner as disclosed above. The steps of the method are preferably carried out with any of the bioactive materials, structures, and base materials disclosed above.

[0086] Of course, polyimide may be deposited as any or all of the porous and additional coating layers 20, 24 and/or 16 by vapor deposition in a manner similar to that disclosed above for parylene and its derivatives. Tech-

niques for the plasma deposition of polymers such as poly(ethylene oxide), poly(ethylene glycol), poly(propylene oxide), silicone, or a polymer of methane, tetrafluoroethylene or tetramethyl-disiloxane on other objects are well-known, and these techniques may be useful in the practice of the present invention.

[0087] In view of the disclosure above, it is clear that the implantable medical device achieves precise control over the release of one or more bioactive materials contained in the device. Moreover, the polyimide, parylene, parylene derivative or other polymeric layers 16, 20 and/or 24 can be remarkably thin, in comparison to the thicknesses required for other polymer layers. The bulk or substantial majority of the overall coating on the structure 12 can therefore consist of bioactive material. This allows the supply of relatively large quantities of bioactive material to the patient, much greater than the amounts supplied by prior devices. These quantities of bioactive material can be supplied to any of a wide variety of locations within a patient during or after the performance of a medical procedure, but are especially useful for preventing abrupt closure and/or restenosis of a blood vessel by the delivery of an antithrombotic or other medication to the region of it which has been opened by PTA. The invention permits the release rate of a bioactive material to be carefully controlled over both the short and long terms. Most importantly, any degradation of the bioactive material which might otherwise occur by other polymer coating techniques is avoided.

[0088] The other details of the construction or composition of the various elements of the disclosed embodiment of the present invention are not believed to be critical to the achievement of the advantages of the present invention, so long as the elements possess the strength or flexibility needed for them to perform as disclosed. The selection of these and other details of construction are believed to be well within the ability of one of ordinary skills in this area, in view of the present disclosure.

Claims

1. An implantable medical device (10), including:

a structure (12) adapted for introduction into a patient, the structure (12) having at least one surface and being composed of a base material (14);

at least one coating layer (16) posited on one surface of the structure (12); and

at least one layer (18) of a bioactive material posited over at least a portion of the at least one coating layer (16), characterised in that the at least one layer (18) of bioactive material forms the outer or outermost layer of the device, and in that release of the bioactive material into a patient is controlled by the said at least one

coating layer (16).

2. The device (10) of claim 1, wherein the at least one coating layer (16) includes a non-porous material or a parylene derivative.

3. The device (10) of claim 2, wherein the at least one coating layer (16) has a thickness in a range from 50,000 to 500,000 Angstroms, more preferably 100,000 to 500,000 Angstroms, or most preferably a thickness of approximately 200,000 Angstroms.

4. The device (10) of claim 2, wherein the non-porous material is at least one of an adsorbent material and an absorbent material having a preferable thickness of approximately 230,000 Angstroms.

5. The device (10) of claim 1, wherein the bioactive material layer (18) includes an antiplatelet GP IIb/IIIa antibody.

6. The device (10) of claim 1, wherein the bioactive material layer (18) includes a chimeric monoclonal antibody such as preferably an antiplatelet GP IIb/IIIa antibody.

7. The device (10) of claim 1, wherein the bioactive material layer (18) includes at least one of: heparin, covalent heparin, or another thrombin inhibitor, hirudin, hirulog, argatroban, D-phenylalanyl-L-poly-L-arginyl chloromethyl ketone, or another antithrombogenic agent, or mixtures thereof; urokinase, streptokinase, a tissue plasminogen activator, or another thrombolytic agent, or mixtures thereof; a fibrinolytic agent; a vasospasm inhibitor; a calcium channel blocker, a nitrate, nitric oxide, a nitric oxide promoter or another vasodilator; an antimicrobial agent or antibiotic; aspirin, ticlopidine, a glycoprotein IIb/IIIa inhibitor or another inhibitor of surface glycoprotein receptors, or another antiplatelet agent; colchicine or another antimitotic, or another microtubule inhibitor, dimethyl sulfoxide (DMSO), a retinoid or another antisecretory agent; cytochalasin or another actin inhibitor; or a remodeling inhibitor; deoxyribonucleic acid, an antisense nucleotide or another agent for molecular genetic intervention; methotrexate or another antimetabolite or antiproliferative agent; tamoxifen citrate, Taxol® or derivatives thereof, or other anti-cancer chemotherapeutic agents; dexamethasone, dexamethasone sodium phosphate, dexamethasone acetate or another dexamethasone derivative, or another anti-inflammatory steroid or non-steroidal anti-inflammatory agent; cyclosporin or another immunosuppressive agent; trapidil (a PDGF antagonist), angiogenin, angiopoietin (a growth hormone antagonist), a growth factor or an anti-growth factor antibody, or another growth factor antagonist; dopamine, bro-

- mocriptine mesylate, pergolide mesylate or another dopamine agonist; ^{60}Co , ^{192}Ir , ^{32}P , ^{111}In , ^{90}Y , $^{99\text{m}}\text{Tc}$ or another radiotherapeutic agent; iodine-containing compounds, barium-containing compounds, gold, tantalum, platinum, tungsten or another heavy metal functioning as a radiopaque agent; a peptide, a protein, an enzyme, an extracellular matrix component, a cellular component or another biologic agent; captopril, enalapril or another angiotensin converting enzyme (ACE) inhibitor; ascorbic acid, alpha tocopherol, superoxide dismutase, deferoxamine, a 21-amino steroid (lasaroid) or another free radical scavenger, iron chelator or antioxidant; a ^{14}C -, ^3H -, ^{131}I -, ^{32}P - or ^{36}S -radiolabelled form or other radiolabelled form of any of the foregoing; estrogen or another sex hormone; AZT or other antipolymerases; acyclovir, famciclovir, rimantadine hydrochloride, ganciclovir sodium or other antiviral agents; 5-aminolevulinic acid, meta-tetrahydroxyphenylchlorin, hexadecafluoro zinc phthalocyanine, tetramethyl hematoporphyrin, rhodamine 123 or other photodynamic therapy agents; an IgG2 Kappa antibody against pseudomonas aeruginosa exotoxin A and reactive with A431 epidermoid carcinoma cells, monoclonal antibody against the noradrenergic enzyme dopamine beta-hydroxylase conjugated to saporin or other antibody targeted therapy agents; gene therapy agents; and enalapril and other prodrugs, or a mixture of any of these.
8. The device (10) of any one of the preceding claims, wherein alternating layers of coating material (20) and bioactive material(s) are applied to the outermost surface of the said bioactive layer (18), the outermost layer being a bioactive layer.
9. The device (10) of claim 1 further including at least one adhesion promotion layer (30) posited on one surface of the structure (12) of which the coating layer (16) is posited over at least a portion of the adhesion promotion layer (30).
- Patentansprüche**
1. Implantierbares medizinisches Gerät (10), aufweisend:
- ein strukturelles Teil (12), das zum Einführen in einen Patienten geeignet ist, wobei das strukturelle Teil (12) mindestens eine Oberfläche aufweist und aus einem Grundmaterial (14) besteht:
- mindestens eine Beschichtungsschicht (16), die auf einer Oberfläche des strukturellen Teils (12) abgesetzt ist; und
- mindestens eine Schicht (18) eines bioaktiven Materials, das auf mindestens einem Teil der mindestens einen Beschichtungsschicht (16) abgesetzt ist, **dadurch gekennzeichnet, dass** mindestens eine Schicht (18) bioaktives Material die äußere oder äußerste Schicht des Geräts bildet und dass die Abgabe des bioaktiven Materials in einen Patienten durch die mindestens eine Beschichtungsschicht (16) gesteuert ist.
2. Gerät (10) nach Anspruch 1, wobei die mindestens eine Beschichtungsschicht (16) ein nichtporöses Material oder ein Parylenderivat aufweist.
3. Gerät (10) nach Anspruch 2, wobei die mindestens eine Beschichtungsschicht (16) eine Dicke im Bereich von 50.000 bis 500.000 Ångström, noch bevorzugter 100.000 bis 500.000 Ångström oder am bevorzugtesten eine Dicke von etwa 200.000 Ångström aufweist.
4. Gerät (10) nach Anspruch 2, wobei das nichtporöse Material mindestens eines unter einem adsorptionsfähigen Material und einem absorptionsfähigen Material mit bevorzugt einer Dicke von etwa 230.000 Ångström ist.
5. Gerät (10) nach Anspruch 1, wobei die Schicht bioaktives Material (18) einen Antithrombozyten-GP IIb/IIIa-Antikörper aufweist.
6. Gerät (10) nach Anspruch 1, wobei die Schicht (18) bioaktives Material einen chimären monoklonalen Antikörper wie bevorzugt einen Antithrombozyten-GP IIb/IIIa-Antikörper aufweist.
7. Gerät (10) nach Anspruch 1, wobei die Schicht (18) bioaktives Material mindestens eines aufweist von: Heparin, kovalentem Heparin oder einem anderen Thrombininhibitor, Hirudin, Hirulog, Argatroban, D-Phenylalanyl-L-poly-L-arginylchloromethylketon oder einem anderen antithrombogenen Mittel oder Mischungen derselben; Urokinase, Streptokinase, einem Gewebeplasminogenaktivator oder einem anderen thrombolytischen Mittel oder Mischungen derselben; einem fibrinolytischen Mittel; einem Vasoconstrictorinhibitor; einem Calciumblocker, einem Nitrat, Stickstoffmonoxid, einem Stickstoffmonoxidpromotor oder einem anderen Vasodilatator; einem antimikrobiellen Mittel oder Antibiotikum; Aspirin, Ticlopidin, einem Glycoprotein IIb/IIIa-Inhibitor oder einem anderen Inhibitor von Oberflächen-Glycoproteinrezeptoren oder einem anderen Antithrombozytenmittel; Colchicin oder einem anderen Anti-

mitotikum oder einem anderen Mikrotubulusinhibitor, Dimethylsulfoxid (DMSO), einem Retinoid oder einem anderen antisekretorischen Mittel; Cytochalasin oder einem anderen Actininhibitor oder einem Remodellierinhibitor; Desoxyribonucleinsäure, einem Antisense nucleotid oder einem anderen Mittel für die molekulare genetische Intervention; Methotrexat oder einem anderen Antimetaboliten oder antiproliferativen Mittel; Tamoxifen citrat, Taxol® oder Derivate desselben oder anderen antineoplastischen chemotherapeutischen Mitteln; Dexamethason, Dexamethasonnatriumphosphat, Dexamethasonacetat oder einem anderen Dexamethasonderivat oder einem anderen entzündungshemmenden Steroid oder nichtsteroidalen entzündungshemmenden Mittel; Cyclosporin oder einem anderen immunsuppressiven Mittel; Trapidal (einem PDGF-Antagonisten), Angiogenin, Angiopoietin (einem Wachstumshormon-Antagonisten), einem Wachstumsfaktor- oder einem Anti-Wachstumsfaktor-Antikörper oder einem anderen Wachstumsfaktorantagonisten; Dopamin, Bromocriptinmesylat, Pergolidmesylat oder einem anderen Dopaminagonisten; ^{60}Co , ^{192}Ir , ^{32}P , ^{111}In , ^{90}Y , $^{99\text{m}}\text{Tc}$ oder einem anderen radiotherapeutischen Mittel, iodhaltigen Verbindungen, bariumhaltigen Verbindungen, Gold, Tantal, Platin, Wolfram oder einem anderen Schwermetall, das als röntgendichtes Mittel wirkt; einem Peptid, einem Protein, einem Enzym, einer extrazellulären Matrixkomponente, einer zellulären Komponente oder einem anderen biologischen Mittel; Captopril, Enalapril oder einem anderen Angiotensin umwandelnden Enzym-(ACE) Inhibitor; Ascorbinsäure, Alpha-Tocopherol, Superoxiddismutase, Deferoxamin, einem 21-AminoSteroid (Lasaroid) oder einem anderen Radikalfänger, Eisenchelator oder Antioxidationsmittel; einer ^{14}C -, ^3H -, ^{131}I -, ^{32}P - oder ^{36}S -radiomarkierten Form oder anderen radiomarkierten Form irgend eines der obigen; Östrogen oder einem anderen Sexualhormon; AZT oder anderen Antipolymerasen; Acyclovir, Famciclovir, Rimantadinhydrochlorid, Ganciclovirnatrium oder anderen antiviralen Mitteln; 5-Aminolevulinsäure, Metatetrahydroxyphenylchlor, Hexadecylfluorizinkphthalocyanin, Tetramethylhämatoporphyrin, Rhodamin 123 oder anderen photodynamischen therapeutischen Mitteln; einem IgG2-Kappa-Antikörper gegen *Pseudomonas aeruginosa*-Exotoxin A, das mit A431-epidermoiden Krebszellen reaktiv ist, monoklonalem Antikörper gegen das noradrenerge Enzym Dopamin-Beta-Hydroxylase, die an Saporin oder anderen auf Antikörper gerichtete therapeutische Mittel konjugiert ist; Gentherapiemitteln; und Enalapril oder anderen Prodrugs oder einer Mischung irgendwelcher derselben.

8. Gerät (10) nach einem der vorhergehenden An-

sprüche, wobei wechselnde Schichten von Beschichtungsmaterial (20) und bioaktivem Material bzw. bioaktiven Materialien auf die äußerste Oberfläche der bioaktiven Schicht (18) aufgebracht werden, wobei die äußerste Schicht eine bioaktive Schicht ist.

9. Gerät (10) nach Anspruch 1, des Weiteren mindestens eine haftungsfördernde Schicht (30) aufweisend, die auf einer Oberfläche des strukturellen Teils (12) aufgebracht ist, bei dem die Beschichtungsschicht (16) über mindestens einen Teil der haftungsfördernden Schicht (30) aufgebracht ist.

Revendications

1. Dispositif médical implantable (10), comprenant :

une structure (12) conçue pour une introduction dans un patient, la structure (12) ayant au moins une surface et étant composée d'un matériau de base (14);

au moins une couche de revêtement (16) appliquée sur une surface de la structure (12); et au moins une couche (18) d'un matériau bioactif appliquée sur au moins une partie de la au moins une couche de revêtement (16), **caractérisé en ce que** la au moins une couche (18) de matériau bioactif forme la couche externe ou le plus à l'extérieur du dispositif et **en ce que** la libération du matériau bioactif dans un patient est réglée par ladite au moins une couche de revêtement (16).

2. Dispositif (10) selon la revendication 1, dans lequel la au moins une couche de revêtement (16) comprend un matériau non poreux ou un dérivé de parylène.

3. Dispositif (10) selon la revendication 2, dans lequel la au moins une couche de revêtement (16) a une épaisseur qui se situe dans une plage de 50 000 à 500 000 angströms, mieux encore dans une plage de 100 000 à 500 000 angströms ou bien mieux encore une épaisseur d'environ 200 000 angströms.

4. Dispositif (10) selon la revendication 2, dans lequel le matériau non poreux est au moins un matériau adsorbant ou un matériau absorbant ayant une épaisseur préférée d'environ 230 000 angströms.

5. Dispositif (10) selon la revendication 1, dans lequel la couche de matériau bioactif (18) comprend un anticorps GP IIb/IIIa antiplaquettaire.

6. Dispositif (10) selon la revendication 1, dans lequel la couche de matériau bioactif (18) comprend un

anticorps monoclonal chimérique, tel que, de préférence, un anticorps GP IIb/IIIa antiplaquettaire.

7. Dispositif (10) selon la revendication 1, dans lequel la couche de matériau bioactif (18) comprend au moins un des matériaux suivants : l'héparine, l'héparine covalente ou un autre inhibiteur de thrombine, l'hirudine, l'hirulog, l'argatroban, la D-phénylalanine-L-poly-L-arginyl chlorométhylcétone ou un autre agent antithrombogénique, ou leurs mélanges; l'urokinase, la streptokinase, un activateur plasminogène de tissu ou un autre agent thrombolytique, ou leurs mélanges; un agent fibrinolytique; un inhibiteur de vasospasme; un agent de blocage du canal de calcium, un nitrate, l'oxyde nitrique, un promoteur d'oxyde nitrique ou un autre vasodilatateur; un agent antimicrobien ou un antibiotique; l'aspirine, la ticlopidine, un inhibiteur IIb/IIIa de glycoprotéine ou un autre inhibiteur de récepteurs de glycoprotéine en surface ou un autre agent antiplaquettaire; la colchicine ou un autre antimitotique ou un autre inhibiteur de microtubule, le diméthylsulfoxyde (DMSO), un rétinoïde ou un autre agent antisécrétoire; la cytochalasine ou un autre inhibiteur d'actine; ou un inhibiteur de remodelage; l'acide désoxyribonucléique, un nucléotide antisense ou un autre agent d'intervention génétique moléculaire; le méthotrexate ou un autre agent antimétabolite ou antiproliférant; le citrate de tamoxifène, le Taxol® ou ses dérivés, ou d'autres agents chimiothérapeutiques anti-cancer; la dexaméthasone, le phosphate sodique de dexaméthasone, l'acétate de dexaméthasone ou un autre dérivé de dexaméthasone, ou un autre stéroïde anti-inflammatoire ou un agent anti-inflammatoire non stéroïdien; la cyclosporine ou un autre agent immunosuppresseur; le trapidal (un antagoniste de PDGF), l'angiogénine, l'angiotensine (un antagoniste de l'hormone de croissance), un anticorps du facteur de croissance ou du facteur d'anti-croissance ou un autre antagoniste du facteur de croissance; la dopamine, le mésylate de bromocriptine, le mésylate de pergolide ou un autre agoniste de la dopamine; ^{60}Co , ^{192}Ir , ^{32}P , ^{111}In , ^{90}Y , $^{99\text{m}}\text{Tc}$ ou un autre agent radiothérapeutique; des composés contenant de l'iode, des composés contenant du baryum, l'or, le tantale, le platine, le tungstène ou un autre métal lourd fonctionnant comme agent opaque aux rayons X; un peptide, une protéine, une enzyme, un composant de matrice extracellulaire, un composant cellulaire ou un autre agent biologique; le captopril, l'énalapril ou un autre inhibiteur d'enzyme de conversion d'angiotensine (ACE); l'acide ascorbique, l'alpha-tocophérol, le peroxyde de dismutase, la déféroxamine, un 21-aminostéroïde (lasaroïde) ou un autre adsorbant de radicaux libres, un chélateur de fer ou un antioxydant; une forme radiomarquée ^{14}C -, ^3H -, ^{131}I -, ^{32}P - ou ^{36}S -

ou une autre forme radiomarquée d'un quelconque des types précités; l'oestrogène ou une autre hormone sexuelle; l'AZT ou d'autres antipolymérase; l'acyclovir, le famciclovir, le chlorhydrate de rimantadine, le ganciclovir sodique ou d'autres agents antiviraux; l'acide 5-aminolévulinique, la méta-tétrahydroxyphénylchlorine, l'hexadécafluorophtalocyanine de zinc, la tétraméthylhématoporphyrine, la rhodamine 123 ou d'autres agents de thérapie photodynamiques; un anticorps Kappa IgG2 contre l'exotoxine A de la souche *Pseudomonas aeruginosa* et réactif avec des cellules du carcinome épidermoïde A431, un anticorps monoclonal contre l'enzyme noradrénergique, la bêta-hydroxylase de dopamine conjuguée à la saporine ou d'autres agents de thérapie ciblés sur les anticorps; les agents de thérapie génique; et l'énalapril et d'autres précurseurs de médicaments, ou un mélange de l'un quelconque de ceux-ci.

8. Dispositif (10) selon l'une quelconque des revendications précédentes, dans lequel des couches alternées de matériau de revêtement (20) et de matériau(x) bioactif(s) sont appliquées à la surface le plus à l'extérieur de ladite couche bioactive (18), la couche le plus à l'extérieur étant une couche bioactive.
9. Dispositif (10) selon la revendication 1, comprenant en outre au moins une couche de promotion d'adhérence (30) appliquée sur une surface de la structure (12) dont la couche de revêtement (16) est appliquée sur au moins une partie de la couche de promotion d'adhérence (30).

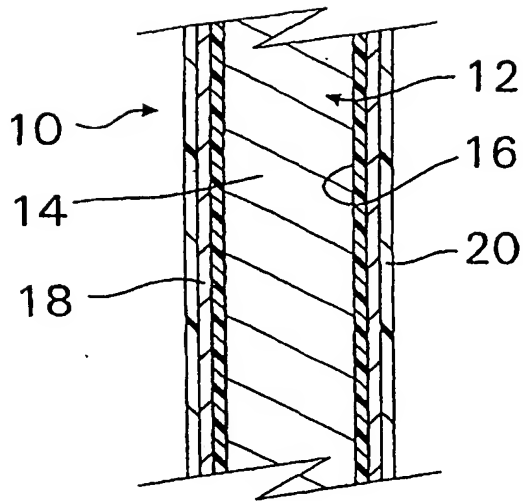


Fig. 1

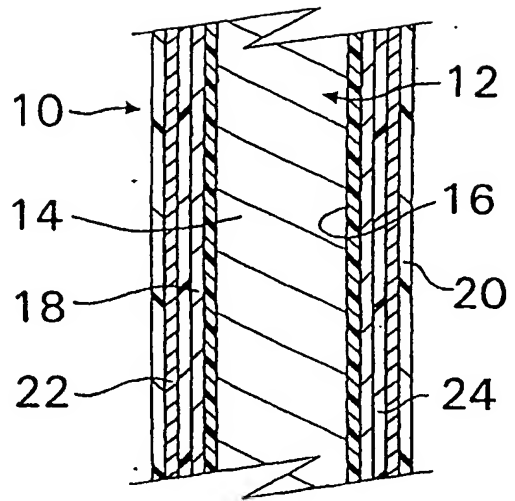


Fig. 2

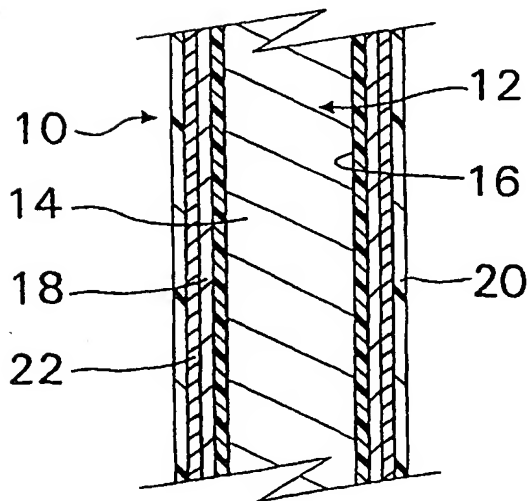


Fig. 3

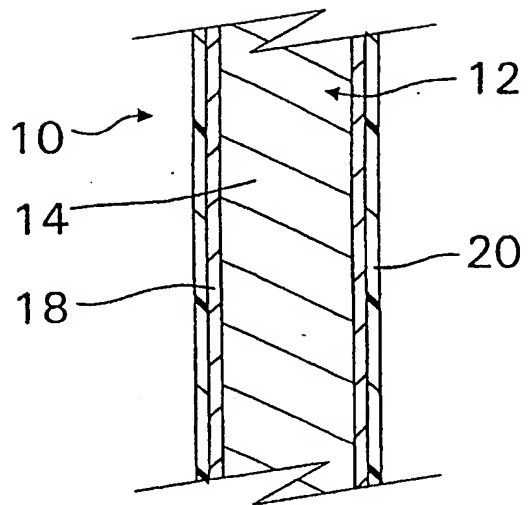


Fig. 4

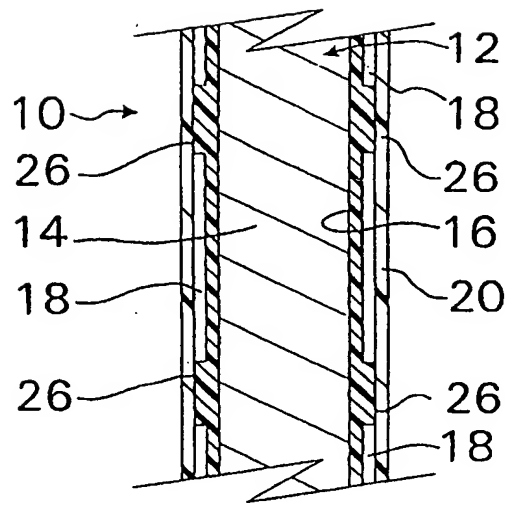


Fig. 5

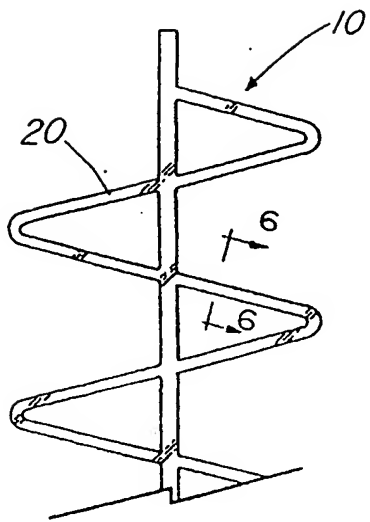


FIG. 7

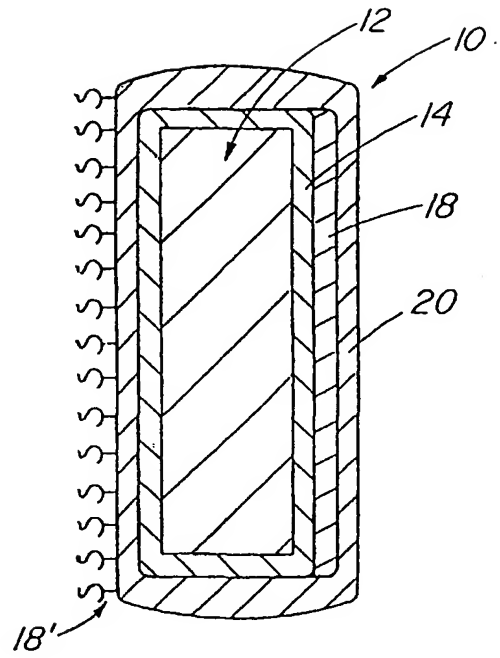


FIG. 6A

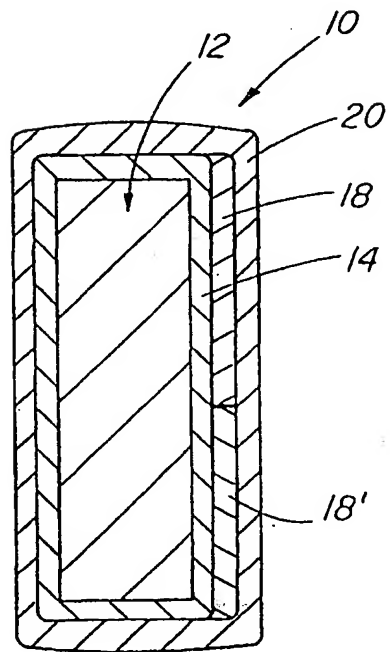
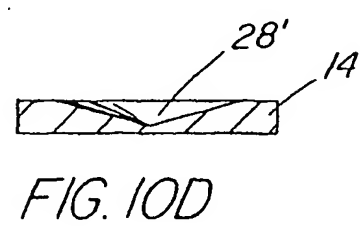
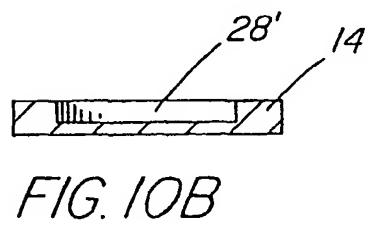
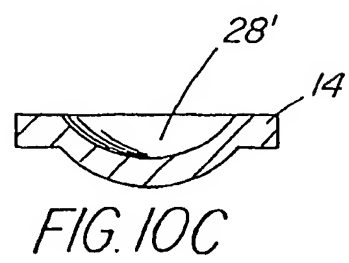
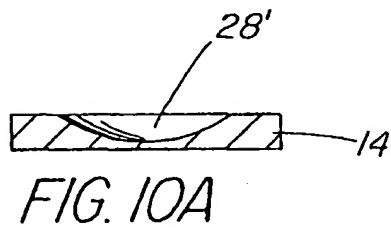
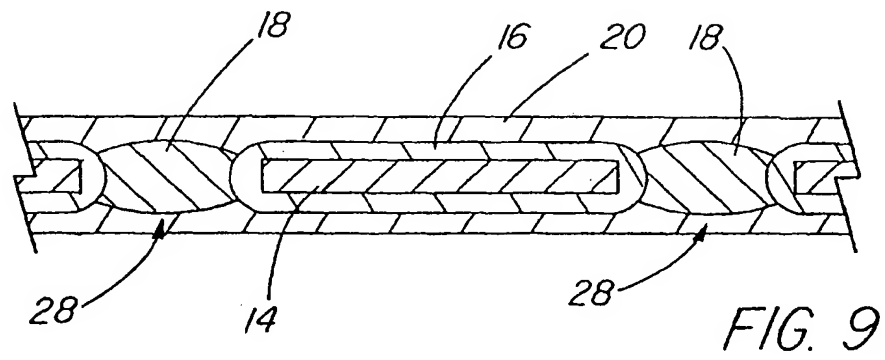
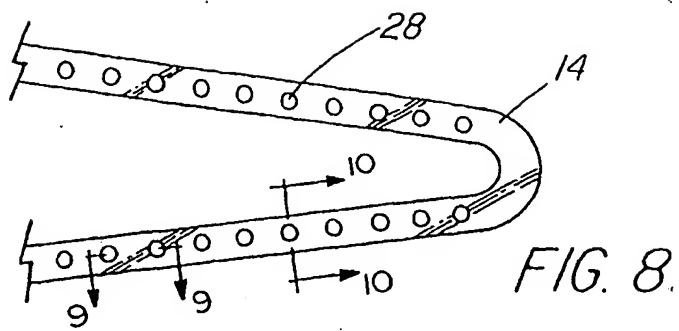


FIG. 6B



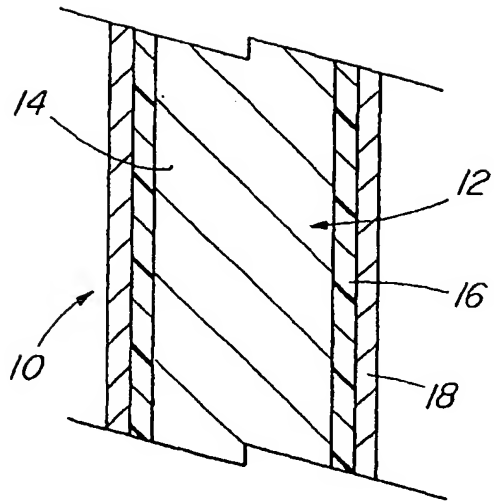


FIG. 11

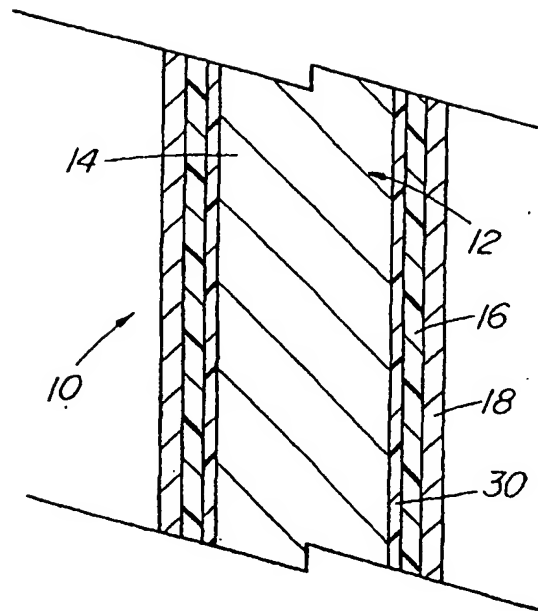


FIG. 12